

All requests for Spravato (esketamine) require a prior authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

Coverage may be provided with a <u>diagnosis</u> of [treatment-resistant depression] and the following criteria is met:

- The member is 18 years of age and older
- Must have a diagnosis of moderate to severe major depressive disorder without psychotic features
- Provider attestation of the following:
 - The diagnosis was made using DSM-5 criteria by or in consultation with a mental health provider
 - Spravato (esketamine) will be used in combination with an oral antidepressant
 - Spravato (esketamine) will be administered under the supervision of a healthcare provider and the member will be monitored for at least 2 hours after administration
 - The member has been assessed and determined not to be at risk for abuse and misuse of Spravato (esketamine)
- If the request is for a new start, documentation showing the member has tried and failed or had an intolerance or contraindication for at least 4 weeks each to at least 2 antidepressants
 - At least one failure must have occurred in the past 3 months
- Documentation of a baseline Montgomery-Åsberg Depression Rating Scale (MADRS) total score.
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- Initial Duration of Approval: 3 months
- Reauthorization criteria
 - Documentation the member responded to therapy demonstrated by an improvement from baseline in MADRS total score
- Reauthorization Duration of Approval: 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

When criteria are not met, the request will be forwarded to a Medical Director for review. The physician reviewer must override criteria when, in their professional judgment, the requested medication is medically necessary.



	SPRAVATO					
Please complete and fax all reques	PRIOR AUTHO			aboratory tast regults, or abort do	aumontation	
	icable to Gateway Health SM Pl				cumentation	
If needed, you may call to speak to a Pharmacy Services Representative.						
	IONE: (800) 392-1147 Monda					
	PROVIDER					
Requesting Provider:			NPI:			
Provider Specialty:			Office Contact:			
Office Address:			Office Phone:			
			Office Fax:			
	MEMBER I	NFORMA	TION			
Member Name: DOB:						
			ember weight: pounds or kg			
REQUESTED DRUG INFORMATION						
Medication: Strength:						
Frequency:			Duration:			
Is the member currently receiving requested medication? Yes			No Date Medication Initiated:			
Billing Information						
This medication will be billed: at a pharmacy OR						
	medically (if medically please	e provide a	JCODE:			
Place of Service: Hospital		per's home				
Place of Service Information						
Name:			NPI:			
Address:			Phone:			
MEDICAL HISTORY (Complete for ALL requests)						
Diagnosis: Treatment Resistant Severe Major Depressive Disorder without psychotic features Other:						
(please provide documentation of how diagnosis was determined)						
Was the diagnosis made using DSM-5 criteria by or in consultation with a mental health provider?						
Please select all of the following that apply:						
The medication will be used in combination with an oral antidepressant						
The medication will be administered under the supervision of a healthcare provider						
The member will be monitored for at least 2 hours after each administration of the medication by a healthcare provider						
The member has been assessed by a health care provider and has been determined not to be at risk for abuse or misuse of the						
requested medication						
Please provide the member's baseline Montgomery-Asberg Depression Rating Scale (MADRS) score:						
CURRENT or PREVIOUS THERAPY						
Medication Name	Strength/ Frequency	Dates of	f Therapy	Status (Discontinued & Why	y/Current)	
				· · · · ·	·	
REAUTHORIZATION						
Has the member experienced a significant improvement with treatment? Yes No						
Please provide the member's curre						
S	UPPORTING INFORMATI	ON or CL	INICAL RA	TIONALE		
Prescribing Provi	der Signature			Date		